

Centre No.	G81688
EQA	2019 Prenatal constitutional CNV detection (previously Prenatal microarray)
Year	2019

Member of the UK NEQAS Consortium

INDIVIDUAL LABORATORY REPORT

To be read in conjunction with the EQA Summary Report

EQA Assessment

Final

A. Case Assessment

Case 1

Category	Score	Comments
Analysis	1.50	The results should be given according to ISCN 2016. (-0.5 marks)
Interpretation	2.00	The written description should include the genetic sex according to the array profile. (-0 marks)
Clerical Accuracy	2.00	The list of genes is unhelpful - only the clinically relevant genes should be discussed. (-0 marks)
		There is no authorisation of the report. (-0 marks)
		The report should be paginated (e.g. page 1 of 1). (-0 marks)

Case 2

Category	Score	Comments
Analysis	1.50	The results should be given according to ISCN 2016. (-0.5 marks)
Interpretation	1.50	The report should request parental bloods to exclude a familial rearrangement. (-0.5 marks)
Clerical Accuracy	2.00	The list of genes is unhelpful - only the clinically relevant genes should be discussed. (-0 marks)
		Clerical errors same as case 1 (marks deducted in case 1). (-0 marks)

Mean Overall Scores

Category	Score
Analysis	1.50
Interpretation	1.75
Clerical Accuracy	2.00

B. Conclusions

Recommendations	None
Performance Statement	Satisfactory

GenQA is operated under the legal entity of OUH NHS Foundation Trust and is provided from two sites.

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Thank you for your participation.

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